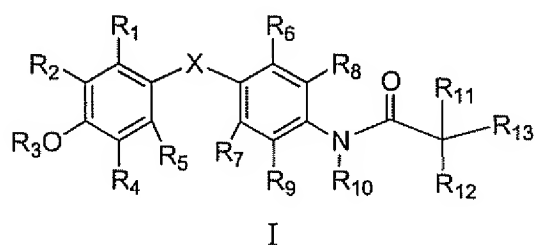


1. Canceled
2. Canceled.
3. (Currently amended) A compound having the following Formula I:



~~The compound as defined in Claim 2~~ wherein

- R<sub>1</sub> is hydrogen;
- R<sub>2</sub> is C<sub>1</sub> to C<sub>6</sub> alkyl or C<sub>3</sub> to C<sub>7</sub> cycloalkyl;
- R<sub>3</sub> is hydrogen;
- R<sub>4</sub> is halogen or C<sub>1</sub> to C<sub>4</sub> alkyl;
- R<sub>5</sub> is hydrogen;
- R<sub>6</sub> and R<sub>7</sub> are independently bromo, chloro or methyl;
- R<sub>8</sub> is halogen or C<sub>1</sub> to C<sub>4</sub> alkyl;
- R<sub>9</sub> is hydrogen or halogen;
- R<sub>10</sub> is hydrogen;
- R<sub>11</sub> is carboxyl;
- R<sub>12</sub> is hydrogen; and
- R<sub>13</sub> is hydrogen.

4. (Original) The compound as defined in Claim 3 wherein R<sub>2</sub> is isopropyl.

5. (Currently amended) The compound as defined in Claim 2 3 wherein

R<sub>1</sub> is hydrogen;

R<sub>2</sub> is isopropyl;

R<sub>3</sub> is hydrogen;

R<sub>4</sub> is C<sub>1</sub> to C<sub>4</sub> alkyl;

R<sub>5</sub> is hydrogen;

R<sub>6</sub> and R<sub>7</sub> are independently bromo, chloro or methyl;

R<sub>8</sub> is halogen or methyl;

R<sub>9</sub> is hydrogen or chloro;

R<sub>10</sub> is hydrogen;

R<sub>11</sub> is carboxyl;

R<sub>12</sub> is hydrogen; and

R<sub>13</sub> is hydrogen.

6. (Currently amended) The compound as defined in Claim 2 3 wherein

R<sub>1</sub> is hydrogen;

R<sub>2</sub> is isopropyl;

R<sub>3</sub> is hydrogen;

R<sub>4</sub> is methyl;

R<sub>5</sub> is hydrogen;

R<sub>6</sub> and R<sub>7</sub> are independently bromo or chloro;

R<sub>8</sub> is chloro or methyl;

R<sub>9</sub> is hydrogen;

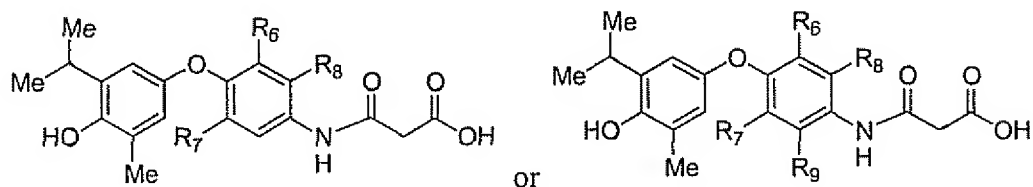
R<sub>10</sub> is hydrogen;

R<sub>11</sub> is carboxyl;

R<sub>12</sub> is hydrogen; and

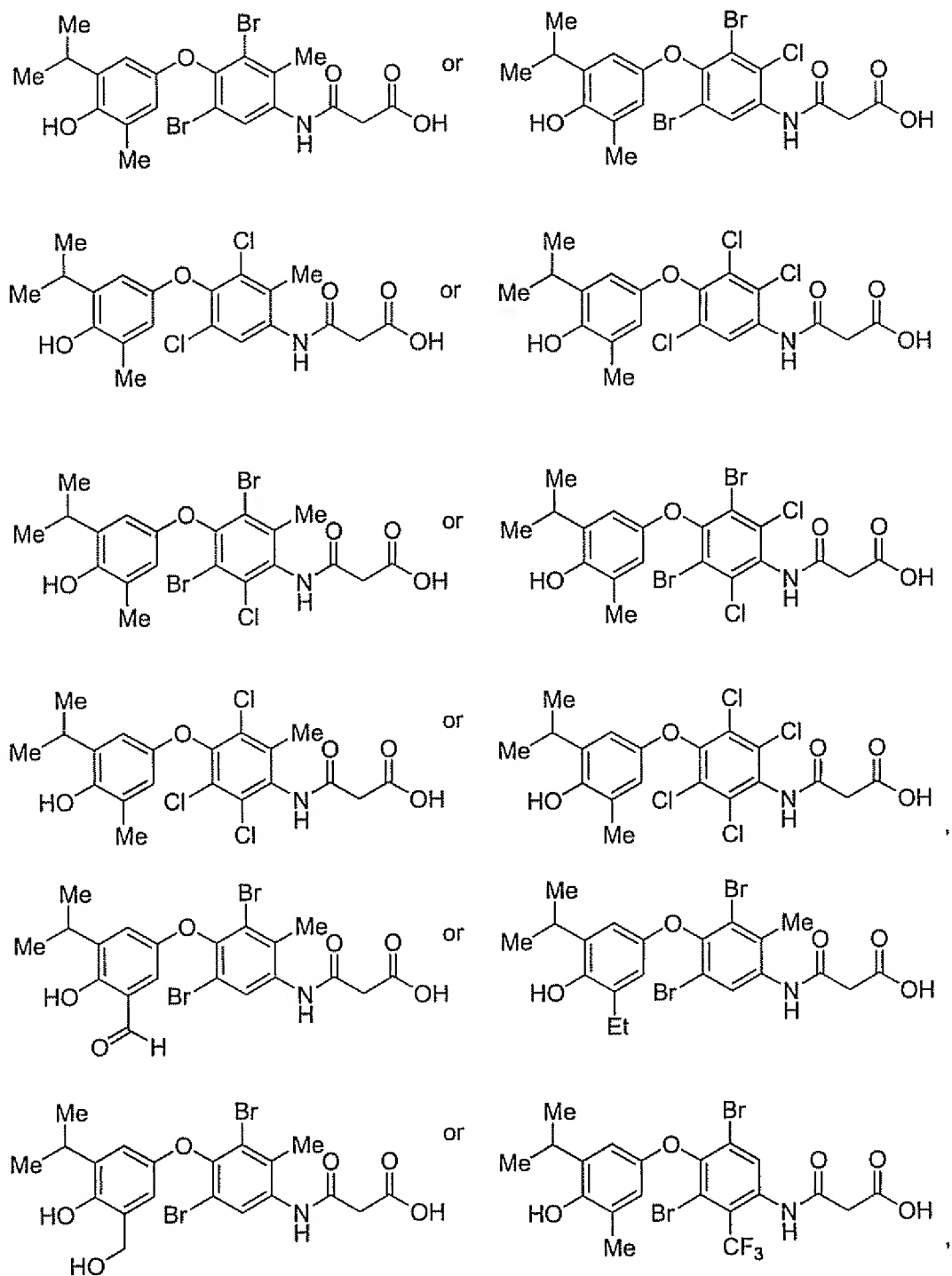
R<sub>13</sub> is hydrogen.

7. (Currently amended) The compound as defined in Claim 4 3 having the structure



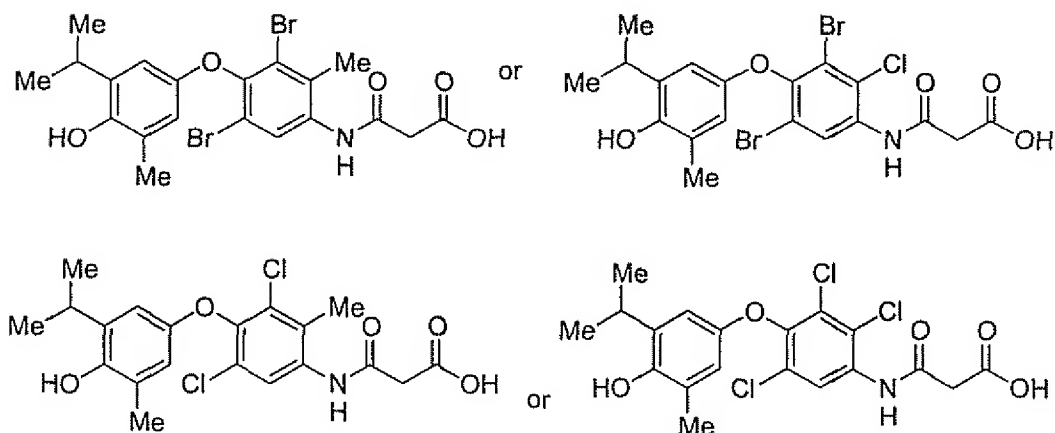
or an alkyl ester thereof.

8. (Currently amended) The compound as defined in Claim 4 3 having the structure

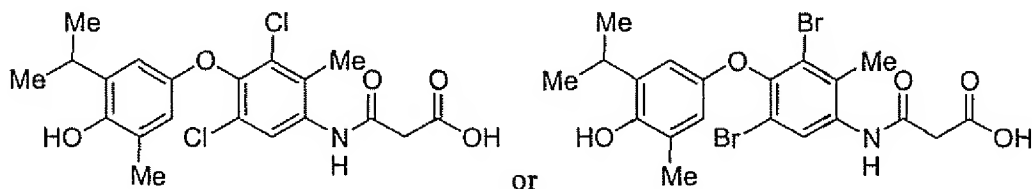


or an alkyl ester thereof.

9. (Currently amended) The compound as defined in Claim 1 3 having the structure



10. (Currently amended) The compound as defined in Claim 1 3 having the structure



11. (Currently amended) A pharmaceutical composition comprising a compound as defined in claim 1 3 and a pharmaceutically acceptable carrier therefor.

12. (Original) The pharmaceutical composition of claim 11 further comprising at least one additional therapeutic agent selected from other compounds of formula I, anti-diabetic agents, anti-osteoporosis agents, anti-obesity agents, growth promoting agents, anti-inflammatory agents, anti-anxiety agents, anti-depressants, anti-hypertensive agents, cardiac glycosides, cholesterol/lipid lowering agents, appetite suppressants, bone resorption inhibitors, thyroid mimetics, anabolic agents, anti-tumor agents and retinoids.

13. (Original) The pharmaceutical composition of claim 12 wherein said additional therapeutic agent is an antidiabetic agent selected from a biguanide, a glucosidase inhibitor, a meglitinide, a sulfonylurea, a thiazolidinedione, a PPAR-alpha agonist, a PPAR-gamma agonist, a PPAR alpha/gamma dual agonist, an SGLT2 inhibitor, a glycogen phosphorylase inhibitor, an  $\alpha$ P2 inhibitor, a glucagon-like peptide-1 (GLP-1), a dipeptidyl peptidase IV inhibitor and insulin.

14. (Original) The pharmaceutical composition of claim 12 wherein said additional therapeutic agent is an antidiabetic agent selected from metformin, glyburide, glimepiride, glipyrider, glipizide, chlorpropamide, gliclazide, acarbose, miglitol, troglitazone, pioglitazone, englitazone, darglitazone, rosiglitazone and insulin.

15. (Original) The pharmaceutical composition of claim 12 wherein said additional therapeutic agent is an anti-obesity agent selected from an  $\alpha$ 2 inhibitor, a PPAR gamma antagonist, a PPAR delta agonist, a beta 3 adrenergic agonist, a lipase inhibitor, a serotonin reuptake inhibitor, a cannabinoid-1 receptor antagonist and an anorectic agent.

16. (Original) The pharmaceutical composition of claim 12 wherein said additional therapeutic agent is a hypolipidemic agent selected from thiazolidinedione, an MTP inhibitor, a squalene synthetase inhibitor, an HMG CoA reductase inhibitor, a fibric acid derivative, an ACAT inhibitor, a cholesterol absorption inhibitor, an ileal  $\text{Na}^+$ /bile cotransporter inhibitor, a bile acid sequestrant and a nicotinic acid or a derivative thereof.

17. (Currently amended) A method for preventing, inhibiting or treating a disease associated with metabolic dysfunction, or which is dependent on the expression of a  $\text{T}_3$  regulated gene, which comprises administering to a mammalian patient in need of treatment a therapeutically effective amount of a compound as defined in claim 4 3.

18. (Currently amended) A method for treating or delaying the progression or onset of obesity, hypercholesterolemia, atherosclerosis, depression, osteoporosis, hypothyroidism, subclinical hyperthyroidism, non-toxic goiter, reduced bone mass, density or growth, eating disorders, reduced cognitive function, thyroid cancer, glaucoma, cardiac arrhythmia, congestive heart failure or a skin disorder or disease, which comprises administering to mammalian patient in need of treatment a therapeutically effective amount of a compound as defined in claim 3 4.

19. (Original) The method according to claim 18 wherein the skin disorder or disease is dermal atrophy, post surgical bruising caused by laser resurfacing, keloids, stria, cellulite, roughened skin, actinic skin damage, lichen planus, ichthyosis, acne, psoriasis, Dernier's disease, eczema, atopic dermatitis, chloracne, pityriasis or skin scarring.

20. (Original) The method according to claim 18 further comprising administering, concurrently or sequentially, a therapeutically effective amount of at least one additional therapeutic agent selected from other compounds of formula I, anti-diabetic agents, anti-osteoporosis agents, anti-obesity agents, growth promoting agents, anti-inflammatory agents, anti-anxiety agents, anti-depressants, anti-hypertensive agents, cardiac glycosides, cholesterol/lipid lowering agents, appetite suppressants, bone resorption inhibitors, thyroid mimetics, anabolic agents, anti-tumor agents and retinoids.

21. (Currently amended) A method of treating or delaying the progression or onset of a skin disorder or disease which comprises administering to a mammalian patient a therapeutically effective amount of a compound as defined in claim 3 ± in combination with a retinoid or a vitamin D analog.

22. (Currently amended) A method for treating or delaying the progression or onset of obesity which comprises administering to mammalian patient in need of treatment a therapeutically effective amount of a compound as defined in Claim 3 ±.

23. (Original) A method according to claim 22 further comprising administering, concurrently or sequentially, a therapeutically effective amount of at least one additional therapeutic agent selected from an anti-obesity agent or an appetite suppressant.

24. (Original) A method according to claim 23 wherein said anti-obesity agent is selected from aP2 inhibitors, PPAR gamma antagonists, PPAR delta agonists, beta 3 adrenergic agonists, lipase inhibitors, serotonin (and dopamine) reuptake inhibitors, cannabinoid-1 receptor antagonists, other thyroid receptor agents and anorectic agents.

25. (Currently amended) A pharmaceutical composition which functions as a selective agonist of the thyroid hormone receptor comprising a compound as defined in claim 3 ±.